

AMENDMENTS TO THE CLAIMS

1. **(Cancelled)**
2. **(Original)** A pharmaceutical composition comprising C-peptide together with at least one pharmaceutically acceptable carrier or excipient for administration to a patient as a once daily dose, wherein said once daily dose does not include a continuous administration or the presence of release rate-controlling agents for the treatment of diabetes and/or microvascular diabetic complications.
3. **(Currently amended)** Method A method of treating diabetes and/or microvascular diabetic complications comprising administering C-peptide or a pharmaceutical composition comprising C-peptide to a patient in a once daily dose, wherein said once daily dose does not include a continuous administration or the presence of release rate-controlling agents.
4. **(Currently amended)** ~~Use,~~ The pharmaceutical composition ~~or method~~ according to Claim 2 ~~any one of claims 1 to 3~~ wherein the C-peptide is human C-peptide
5. **(Currently amended)** ~~Use,~~ The pharmaceutical composition ~~or method~~ according to Claim 2 ~~any one of claims 1 to 4~~ wherein said C-peptide is a ~~the~~ fragment EGSLQ (SEQ ID NO: [[-]] 2).
6. **(Currently amended)** ~~Use,~~ The pharmaceutical composition ~~or method~~ according to Claim 2 ~~any one of claims 1 to 5~~ wherein the patient is a human.
7. **(Currently amended)** ~~Use,~~ The pharmaceutical composition ~~or method~~ according to Claim 2 ~~any one of claims 1 to 6~~ wherein the medicament contains 100 to 1800 nmol of C-peptide.
8. **(Currently amended)** ~~Use,~~ The pharmaceutical composition ~~or method~~ according to Claim 2 ~~any one of claims 1 to 7~~ wherein the medicament is an uncompromised aqueous solution.
9. **(Currently amended)** ~~Use,~~ The pharmaceutical composition ~~or method~~ according to Claim 2 ~~any one of claims 1 to 8~~ wherein said complications are diabetic nephropathy, retinopathy or neuropathy.
10. **(New)** The method according to Claim 3, wherein the C-peptide is human C-peptide

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11. (New) The method according to Claim 3, wherein said C-peptide is a fragment EGSLQ (SEQ ID NO: 2).

12. (New) The method according to Claim 3, wherein the patient is a human.

13. (New) The method according to Claim 3, wherein the medicament contains 100 to 1800 nmol of C- peptide.

14. (New) The method according to Claim 3, wherein the medicament is an uncompromised aqueous solution.

15. (New) The method according to Claim 3, wherein said complications are diabetic nephropathy, retinopathy or neuropathy.